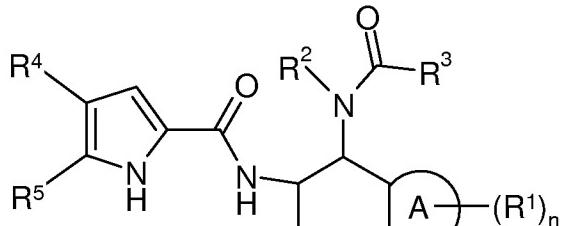


**Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

1. (previously presented) A compound of formula (1):



(1)

wherein:

R<sup>4</sup> and R<sup>5</sup> together are either -S-C(R<sup>6</sup>)=C(R<sup>7</sup>)- or -C(R<sup>7</sup>)=C(R<sup>6</sup>)-S- ;

R<sup>6</sup> and R<sup>7</sup> are independently selected from hydrogen, halo, nitro, cyano, hydroxy, fluoromethyl, difluoromethyl, trifluoromethyl, trifluoromethoxy, carboxy, carbamoyl, (1-4C)alkyl, (2-4C)alkenyl, (2-4C)alkynyl, (1-4C)alkoxy and (1-4C)alkanoyl;

A is phenylene;

n is 0, 1 or 2;

R<sup>1</sup> is independently selected from halo, nitro, cyano, hydroxy, carboxy, carbamoyl, N-(1-4C)alkylcarbamoyl, N,N-((1-4C)alkyl)<sub>2</sub>carbamoyl, sulphamoyl, N-(1-4C)alkylsulphamoyl, N,N-((1-4C)alkyl)<sub>2</sub>sulphamoyl, -S(O)<sub>b</sub>(1-4C)alkyl (wherein b is 0,1,or 2), -OS(O)<sub>2</sub>(1-4C)alkyl, (1-4C)alkyl, (2-4C)alkenyl, (2-4C)alkynyl, (1-4C)alkoxy, (1-4C)alkanoyl, (1-4C)alkanoyloxy, hydroxy(1-4C)alkyl, fluoromethyl, difluoromethyl, trifluoromethyl, trifluoromethoxy and -NHSO<sub>2</sub>(1-4C)alkyl;

or, when n is 2, the two R<sup>1</sup> groups, together with the carbon atoms of A to which they are attached, may form a 4 to 7 membered saturated ring optionally being substituted by one or two methyl groups;

one of R<sup>2</sup> and R<sup>3</sup> is selected from R<sub>Na</sub>, and the other is selected from R<sub>Nb</sub>;

R<sub>Na</sub>: (1-3C)alkyl, halo(1-3C)alkyl, dihalo(1-3)alkyl, trifluoromethyl, hydroxy(1-3C)alkyl, dihydroxy(2-3C)alkyl, cyano(1-3C)alkyl (optionally substituted on alkyl with hydroxy), methoxymethyl, ethoxymethyl, methoxyethyl, methoxymethoxymethyl, dimethoxyethyl, (hydroxy)(methoxy)ethyl, (amino)(hydroxy)(2-3C)alkyl, (aminocarbonyl)(hydroxy)(2-3C)alkyl, (methylaminocarbonyl)(hydroxy)(2-3C)alkyl, (dimethylaminocarbonyl)(hydroxy)(2-3C)alkyl,

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(methylcarbonylamino)(hydroxy)(2-3C)alkyl, (methylS(O)<sub>p</sub>-)(hydroxy)(2-3C)alkyl (wherein p is 0, 1 or 2);

R<sub>Nb</sub>: (1-4C)alkyl, halo(1-4C)alkyl, dihalo(1-4C)alkyl, trifluoromethyl, hydroxy(1-4C)alkyl, dihydroxy(2-4C)alkyl, trihydroxy(3-4C)alkyl, cyano(1-4C)alkyl (optionally substituted on alkyl with hydroxy), (1-4C)alkoxy(1-4C)alkyl, (1-4C)alkoxy(1-4C)alkoxy(1-4C)alkyl, di[(1-4C)alkoxy](2-4C)alkyl, (hydroxy)[(1-4C)alkoxy](2-4C)alkyl, (amino)(hydroxy)(2-4C)alkyl, (aminocarbonyl)(hydroxy)(2-4C)alkyl, ((1-4C)alkylaminocarbonyl)(hydroxy)(2-4C)alkyl, (di(1-4C)alkylaminocarbonyl)(hydroxy)(2-4C)alkyl, ((1-4C)alkylcarbonylamino)(hydroxy)(2-4C)alkyl, ((1-4C)alkylS(O)<sub>p</sub>)(hydroxy)(2-4C)alkyl (wherein p is 0, 1 or 2);

wherein any alkyl or alkoxy group within any group in R<sub>NA</sub> and R<sub>NB</sub> may also optionally be substituted on an available carbon atom with a hydroxy group (provided that said carbon atom is not already substituted by a group linked by a heteroatom); provided that if R<sup>2</sup> is (1-3C)alkyl or (1-4C)alkyl then R<sup>3</sup> is not (1-4C)alkyl or (1-3C)alkyl; or a pharmaceutically acceptable salt thereof.

2. (previously presented) A compound of formula (1) as claimed in Claim 1, or a pharmaceutically acceptable salt thereof, wherein R<sup>2</sup> is selected from R<sub>Na</sub>, and R<sup>3</sup> is selected from R<sub>Nb</sub>, wherein R<sub>Na</sub> and R<sub>Nb</sub> are as defined in Claim 1.

3. (cancelled)

4. (previously presented) A compound of formula (1) as claimed in Claim 1, or a pharmaceutically acceptable salt thereof, wherein n is 0.

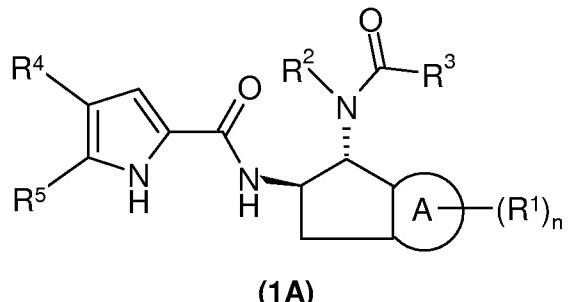
5. (previously presented) A compound of formula (1) as claimed in Claim 1, or a pharmaceutically acceptable salt thereof, wherein R<sup>6</sup> and R<sup>7</sup> are independently selected from hydrogen and halo.

6. (previously presented) A compound of formula (1) as claimed in Claim 1, or a pharmaceutically acceptable salt thereof, wherein R<sup>6</sup> and R<sup>7</sup> are independently selected from hydrogen and chloro.

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7. (previously presented) A compound of formula (1) as claimed in Claim 1, or a pharmaceutically acceptable salt thereof, wherein R<sub>Na</sub> is selected from (1-4C)alkyl, hydroxy(1-4C)alkyl, and (1-4C)alkoxy(1-4C)alkyl.

8. (previously presented) A compound of formula (1) as claimed in Claim 1, or a pharmaceutically acceptable salt thereof, which is a compound of formula (1A):



wherein R<sup>1</sup> to R<sup>7</sup>, A and n are as defined in claim 1.

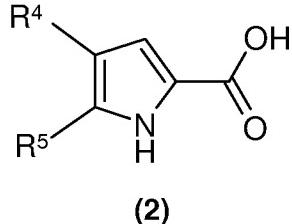
9. (cancelled)

10. (original) A pharmaceutical composition which comprises a compound of the formula (1), or a pharmaceutically acceptable salt thereof, as claimed in claim 1 in association with a pharmaceutically-acceptable diluent or carrier.

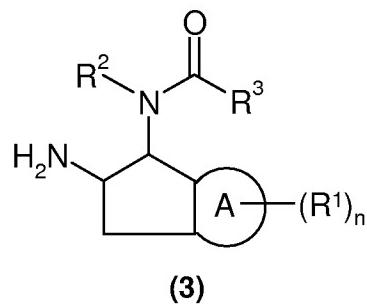
11-15. (cancelled)

16. (currently amended) A process for the preparation of a compound of formula (1) as claimed in claim 1, which process comprises:

reacting an acid of the formula (2):



or an acid chloride or ester activated derivative thereof; with an amine of formula (3):



and thereafter if necessary:

- i) converting a compound of the formula (1) into another compound of the formula (1);
- ii) removing any protecting groups;
- iii) forming a pharmaceutically acceptable salt.

17. (previously presented) A compound of formula (1), or a pharmaceutically acceptable salt thereof, selected from:

2-chloro-N-{(1*R*,2*R*)-1-[(methoxyacetyl)(methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;  
2-chloro-N-{(1*R*,2*R*)-1-[[3-hydroxy-2-(hydroxymethyl)propanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;  
ethyl 3-[(1*R*,2*R*)-2-[(2-chloro-6*H*-thieno[2,3-*b*]pyrrol-5-yl)carbonyl]amino]-2,3-dihydro-1*H*-inden-1-yl](methyl)amino]-3-oxopropanoate;  
2-[(1*R*,2*R*)-2-[(2-chloro-6*H*-thieno[2,3-*b*]pyrrol-5-yl)carbonyl]amino]-2,3-dihydro-1*H*-inden-1-yl](methyl)amino]-2-oxoethyl acetate;  
2-chloro-N-{(1*R*,2*R*)-1-[glycoloyl(methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;  
2-chloro-N-{(1*R*,2*R*)-1-[glyceroyl(methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;  
2-chloro-N-{(1*R*,2*R*)-1-[(2*S*)-2,3-dihydroxypropanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;  
2-chloro-N-{(1*R*,2*R*)-1-[(2*R*)-2,3-dihydroxypropanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;  
2-chloro-N-{(1*R*,2*R*)-1-[(3-hydroxypropanoyl)(methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;  
2-chloro-N-{(1*R*,2*R*)-1-[glycoloyl(2-hydroxyethyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;

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2-chloro-*N*-{(1*R*,2*R*)-1-[(2*R*)-2-hydroxypropanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;  
2-chloro-*N*-{(1*R*,2*R*)-1-[(2*S*)-2-hydroxypropanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;  
2,3-dichloro-*N*-{(1*R*,2*R*)-1-[(2*R*)-2,3-dihydroxypropanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-4*H*-thieno[3,2-*b*]pyrrole-5-carboxamide  
2,3-dichloro-*N*-{(1*R*,2*R*)-1-[(2*S*)-2,3-dihydroxypropanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-4*H*-thieno[3,2-*b*]pyrrole-5-carboxamide;  
(2*S*)-*N*<sup>l</sup>-((1*R*,2*R*)-2-{[(2-chloro-6*H*-thieno[2,3-*b*]pyrrol-5-yl)carbonyl]amino}-2,3-dihydro-1*H*-inden-1-yl)-2-hydroxy-*N*<sup>l</sup>-methylsuccinamide;  
(2*S*)-*N*<sup>l</sup>-((1*R*,2*R*)-2-{[(2,3-dichloro-4*H*-thieno[3,2-*b*]pyrrol-5-yl)carbonyl]amino}-2,3-dihydro-1*H*-inden-1-yl)-2-hydroxy-*N*<sup>l</sup>-methylsuccinamide;  
2,3-dichloro-*N*-{(1*R*,2*R*)-1-[(2*S*)-2-hydroxybutanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-4*H*-thieno[3,2-*b*]pyrrole-5-carboxamide;  
2,3-dichloro-*N*-{(1*R*,2*R*)-1-[(2*S*)-2-hydroxy-3-methylbutanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-4*H*-thieno[3,2-*b*]pyrrole-5-carboxamide;  
2,3-dichloro-*N*-{(1*R*,2*R*)-1-[(2*S*)-4-(1,3-dioxo-1,3-dihydro-2*H*-isoindol-2-yl)-2-hydroxybutanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-4*H*-thieno[3,2-*b*]pyrrole-5-carboxamide;  
2,3-dichloro-*N*-{(1*R*,2*R*)-1-[(2*R*)-2-hydroxy-3-(methylthio)propanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-4*H*-thieno[3,2-*b*]pyrrole-5-carboxamide;  
*tert*-butyl {(2*S*)-3-[(1*R*,2*R*)-2-{[(2,3-dichloro-4*H*-thieno[3,2-*b*]pyrrol-5-yl)carbonyl]amino}-2,3-dihydro-1*H*-inden-1-yl](methyl)amino]-2-hydroxy-3-oxopropyl}carbamate;  
2,3-dichloro-*N*-{(1*R*,2*R*)-1-[(2*S*)-3-cyano-2-hydroxypropanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-4*H*-thieno[3,2-*b*]pyrrole-5-carboxamide;  
*N*-{(1*R*,2*R*)-1-[(*N*-acetylseryl)(methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-2-chloro-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;  
*N*-{(1*R*,2*R*)-1-[(*N*-acetylseryl)(methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-2,3-dichloro-4*H*-thieno[3,2-*b*]pyrrole-5-carboxamide;  
2,3-dichloro-*N*-{(1*R*,2*R*)-1-[methyl(L-seryl)amino]-2,3-dihydro-1*H*-inden-2-yl}-4*H*-thieno[3,2-*b*]pyrrole-5-carboxamide hydrochloride;  
2-chloro-*N*-{(1*R*,2*R*)-1-[methyl(L-seryl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide hydrochloride;  
(2*S*)-*N*<sup>l</sup>-((1*R*,2*R*)-2-{[(2-chloro-6*H*-thieno[2,3-*b*]pyrrol-5-yl)carbonyl]amino}-2,3-dihydro-1*H*-inden-1-yl)-2-hydroxy-*N*<sup>l</sup>-methylpentanediamide;

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(2*S*)-*N*<sup>1</sup>-((1*R,2R*)-2-{[(2,3-dichloro-4*H*-thieno[3,2-*b*]pyrrol-5-yl)carbonyl]amino}-2,3-dihydro-1*H*-inden-1-yl)-2-hydroxy-*N*<sup>1</sup>-methylpentanediamide;  
2-chloro-*N*-{(1*R,2R*)-1-[(2*S*)-2-hydroxy-3-methoxypropanoyl](methyl) amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;  
2,3-dichloro-*N*-{(1*R,2R*)-1-[(2*S*)-2-hydroxy-3-methoxypropanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-4*H*-thieno[3,2-*b*]pyrrole-5-carboxamide;  
2,3-dichloro-*N*-{(1*R,2R*)-1-[(2*R*)-2-hydroxy-3-(methylsulfonyl)propanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-4*H*-thieno[3,2-*b*]pyrrole-5-carboxamide;  
*N*-{(1*R,2R*)-1-[(2*S*)-3-amino-2-hydroxypropanoyl](methyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-2,3-dichloro-4*H*-thieno[3,2-*b*]pyrrole-5-carboxamide hydrochloride;  
(2*S*)-*N*<sup>1</sup>-((1*R,2R*)-2-{[(2,3-dichloro-4*H*-thieno[3,2-*b*]pyrrol-5-yl)carbonyl]amino}-2,3-dihydro-1*H*-inden-1-yl)-2-hydroxy-*N*<sup>1</sup>,*N*<sup>4</sup>-dimethylsuccinamide;  
(2*S*)-*N*<sup>1</sup>-((1*R,2R*)-2-{[(2,3-dichloro-4*H*-thieno[3,2-*b*]pyrrol-5-yl)carbonyl] amino}-2,3-dihydro-1*H*-inden-1-yl)-2-hydroxy-*N*<sup>1</sup>,*N*<sup>4</sup>,*N*<sup>4</sup>-trimethylsuccinamide;  
2-chloro-*N*-{(1*R,2R*)-1-[glyceroyl(2-hydroxyethyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide;  
2-chloro-*N*-{(1*R,2R*)-1-[(2*R*)-2,3-dihydroxypropanoyl](2-hydroxyethyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide; and  
2-chloro-*N*-{(1*R,2R*)-1-[(2*S*)-2,3-dihydroxypropanoyl](2-hydroxyethyl)amino]-2,3-dihydro-1*H*-inden-2-yl}-6*H*-thieno[2,3-*b*]pyrrole-5-carboxamide.

18. (original) A method of producing a glycogen phosphorylase inhibitory effect in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (1) as claimed in claim 1.

19. (original) A method of treating type 2 diabetes, insulin resistance, syndrome X, hyperinsulinaemia, hyperglucagonaemia, cardiac ischaemia or obesity in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (1) as claimed in claim 1.

20. (original) A method of treating type 2 diabetes in a warm-blooded animal, such as man, in need of such treatment which comprises administering to said animal an effective amount of a compound of formula (1) as claimed in claim 1.